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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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09/825,423

04/03/2001

Patricia C. Weber

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2057

24265

7590

10/03/2006

SCHERING-PLOUGH CORPORATION
PATENT DEPARTMENT (K-6-1, 1990)
2000 GALLOPING HILL ROAD
KENILWORTH, NJ 07033-0530

EXAMINER

STEADMAN, DAVID J

ART UNIT

PAPER NUMBER

1656

DATE MAILED: 10/03/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No.

09/825,423

Applicant(s)

WEBER ET AL.

Examiner

David J. Steadman

Art Unit

1656

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --
Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 17 July 2006.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-3,7-9,11,21 and 22 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☒ Claim(s) 1-3,7 and 8 is/are allowed.
- 6) ☒ Claim(s) 9,11,21 and 22 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
 - ☐ Certified copies of the priority documents have been received in Application No. _____.
 - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- ☒ Notice of References Cited (PTO-892)
- ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- ☐ Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date _____
- ☐ Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____
- ☐ Notice of Informal Patent Application
- ☒ Other: Appendices A, B, C.

DETAILED ACTION

Application Status

1. Claims 1-3, 7-9, 11, and 21-22 are pending in the application.
2. Applicant's amendment to the claims, filed on 17 July 2006, is acknowledged.

This listing of the claims replaces all prior versions and listings of the claims.

3. Applicant's amendment to the specification, filed on 17 July 2006, is acknowledged. In view of this statement, sequence compliance appears to be perfected.

4. Applicant's arguments filed on 17 July 2006 in response to the Office action mailed on 7 March 2006 have been fully considered and are deemed to be persuasive to overcome at least one of the rejections and/or objections previously applied. Rejections and/or objections not reiterated from previous office actions are hereby withdrawn.

5. The text of those sections of Title 35, U.S. Code not included in the instant action can be found in a prior Office action.

Claim Rejections - 35 USC § 112, Second Paragraph

6. Claims 9 and 21-22 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claims 9 and 21 are confusing in the recitation of "[a]n isolated polypeptide defined by a variant...the variant consists of a single amino acid substitution..."

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According to MPEP 2111.03, "[t]he transitional phrase 'consisting of' excludes any element, step, or ingredient not specified in the claim." Thus, claims 9 and 21 would appear to read on a polypeptide variant consisting of a single amino acid as defined by the claims. Claim 22 is also rejected as being confusing in the use of the transitional phrase "consists of" in the recitation of "[a]n isolated polypeptide defined by a variant" of SEQ ID NO:5 "wherein the variant consists of a substitution of the amino acids at positions 255-258." In the interest of advancing prosecution, claims 9 and 21 have been interpreted as meaning a variant of SEQ ID NO:3, 5, or 6 with a single amino acid mutation, wherein the mutation is at position 73 or 81. Claim 22 has been interpreted as meaning the polypeptide of SEQ ID NO:5, wherein amino acids 255-258 of SEQ ID NO:5 are replaced with SEQ ID NO:7, 8, 9, 10, 11, 12, 13, or 14. It is suggested that applicant clarify the meaning of the claims.

Claim Rejections - 35 USC § 112, First Paragraph

7. Claim 22 is rejected under 35 U.S.C. § 112, first paragraph, written description, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. This is a new matter rejection.

New claim 22 is drawn to a polypeptide "defined by a variant of...SEQ ID NO:5, wherein the variant consists of a substitution...at positions 255-258 with SEQ ID NO:7,8,9,10,11,12,13, or 14. MPEP § 2163 states, "when filing an amendment an

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applicant should show support in the original disclosure for new or amended claims” and “[i]f the originally filed disclosure does not provide support for each claim limitation, or if an element which applicant describes as essential or critical is not claimed, a new or amended claim must be rejected under 35 U.S.C. 112, para. 1, as lacking adequate written description.” Applicant points to original claim 7, p. 11 of the specification, and “by comparing the full-length sequence in SEQ ID NO:1 with the subdomain I,II fragment sequence in SEQ ID NO:5” (instant response at p. 5, bottom). The examiner has reviewed applicant’s cited supporting disclosure and has aligned SEQ ID NO:1 against SEQ ID NO:5 (see Appendix C). However, this disclosure does not appear to support the claimed variant polypeptide. It is suggested that applicant show support for new claim 22. If applicant maintains that the cited disclosure supports claim 22 as written, applicant is requested to provide a detailed explanation as to how this cited disclosure supports the polypeptide of claim 22.

8. The written description rejection of claim 11 under 35 U.S.C. § 112, first paragraph, is maintained for the reasons of record and the reasons stated below. The rejection was fully explained in the prior Office action.

RESPONSE TO ARGUMENT: Applicant argues the rejection is overcome by claim amendment to “define the polypeptide in the crystalline composition by both a specific amino acid sequence and a specific set of structural coordinates.”

Applicant’s argument is not found persuasive. The examiner maintains the position that the specification fails to describe the genus of crystalline compositions of

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claim 11. While the amendment to the claims limits the *polypeptide* of the composition as the recitation of "crystalline composition" in claim 11 does not specifically define any of the crystalline compositions that fall within its definition, particularly as the recitation of "crystalline composition" does not define any structural features commonly possessed by members of the genus of crystalline compositions of SEQ ID NO:17 that distinguish them from others. One skilled in the art therefore cannot, as one can do with a fully described genus, visualize or recognize the identity of the members of the genus of proteins in crystalline form. In this case, the structure(s) of the genus of *crystals* of the protein of SEQ ID NO:17 is completely undefined.

Applicant appears to take the position that by virtue of limiting the polypeptide of the crystalline composition to SEQ ID NO:17 having the structural coordinates of Table 5, the genus of crystals is adequately described, however, it is well-known in the art that a single polypeptide can crystallize into a plurality of distinct crystal forms, which one cannot predict *a priori* (see, e.g., Aleshin et al. *FEBS Lett* 434:42-46, 1998). Thus, as noted in the prior Office action, the genus of crystals encompasses species that are widely variant, encompassing crystals of unliganded and liganded forms of SEQ ID NO:17, wherein the liganded form is in complex with *any* ligand(s). In this case, the specification discloses only a single representative species of the genus of recited crystalline compositions, *i.e.*, a protein crystal of SEQ ID NO:17 having space group P2₁ and unit cell dimensions $a=34.8 \text{ \AA}$, $b=67.1 \text{ \AA}$, $c=58.4 \text{ \AA}$, $\alpha=\gamma=90^\circ$, and $\beta=101.3^\circ$ (see particularly pp. 41-42 of the specification, which teaches crystallization of SEQ ID NO:17, which is amino acids 181-324 of the HCV NS3 helicase of SEQ ID NO:1), which

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is undisputed by applicant. Other than these single species, the specification fails to describe any other crystals of SEQ ID NO:17 as encompassed by the claims. MPEP § 2163 states “[f]or inventions in an unpredictable art, adequate written description of a genus which embraces widely variant species cannot be achieved by disclosing only one species within the genus.” As such, the single disclosed species of crystals of SEQ ID NO:17 fails to describe all crystals as encompassed by the claim.

Given the lack of description of a representative number of protein crystals, the specification fails to sufficiently describe the claimed invention in such full, clear, concise, and exact terms that a skilled artisan would recognize that applicant was in possession of the claimed invention.

9. The scope of enablement rejection of claim 11 under 35 U.S.C. 112, first paragraph, is maintained for the reasons of record and the reasons stated below. The rejection was fully explained in the prior Office action.

RESPONSE TO ARGUMENT: Applicant argues the rejection is overcome by claim amendment to “define the polypeptide in the crystalline composition by both a specific amino acid sequence and a specific set of structural coordinates.”

Applicant's argument is not found persuasive. The examiner maintains the position that the specification fails to enable all crystals as broadly encompassed by the claim. While the examiner acknowledges the amendment to limit the polypeptide of the crystal to SEQ ID NO:17 having the structural coordinates of Table 5, claim 11 nonetheless broadly encompasses all crystals of SEQ ID NO:17, unliganded or

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complexed with any ligand, having any space group, and any unit cell dimensions. The specification discloses only a single working example of the claimed crystal, *i.e.*, a crystal of SEQ ID NO:17 having space group $P2_1$ and unit cell dimensions $a=34.8 \text{ \AA}$, $b=67.1 \text{ \AA}$, $c=58.4 \text{ \AA}$, $\alpha=\gamma=90^\circ$, and $\beta=101.3^\circ$ (see particularly pp. 41-42 of the specification, which teaches crystallization of SEQ ID NO:17, which is amino acids 181-324 of the HCV NS3 helicase of SEQ ID NO:1). The specification fails to disclose any other working examples or guidance for making other protein crystals of SEQ ID NO:17 under any other conditions with an expectation of obtaining diffraction-quality crystals. As noted in the prior Office action – and undisputed by application – the state of the art at the time of the invention acknowledges a high level of unpredictability for making a protein crystal. For example, the reference of Branden et al. (“Introduction to Protein Structure Second Edition”, Garland Publishing Inc., New York, 1999; cited in the prior Office action) teaches that “[c]rystallization is usually quite difficult to achieve” (p. 375) and that “[w]ell-ordered crystals...are difficult to grow because globular protein molecules are large, spherical, or ellipsoidal objects with irregular surfaces, and it is impossible to pack them into a crystal without forming large holes or channels between the individual molecules” (p. 374). Also, Drenth et al. (“Principles of X-ray Crystallography,” Springer, New York, 1995; cited in the prior Office action) teaches that “[t]he science of protein crystallization is an underdeveloped area” and “[p]rotein crystallization is mainly a trial-and-error procedure” (p. 1). One cannot predict *a priori* those conditions that will lead to the successful crystallization of a diffraction-quality crystal nor can one predict the space group symmetry or unit cell dimensions of the

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resulting crystal. As stated above, even a single polypeptide can have multiple crystal forms, however, what form will result from which particular crystallization conditions – if any – remains highly unpredictable as evidenced by the state of the art at the time of the invention. While applicant may argue that a crystal of SEQ ID NO:17 in complex with a ligand or ligands can be prepared according to the disclosed method and would have the same space group and unit cell dimensions, there is no way to predict *a priori* the space group and unit cell dimensions of a protein, as evidenced by the references of Kierzek et al. (cited in the prior Office action; see cited relevant teachings). While methods of protein crystallography were known at the time of the invention, it was not routine in the art to make all polypeptide crystals as encompassed by the claims and screen for those that are diffraction-quality under any crystallization conditions as encompassed by the claims, diffract those crystals, and to determine those polypeptide crystal structures that represent biologically-relevant macromolecules.

In view of the overly broad scope of the claims, the lack of guidance and working examples provided in the specification, the high level of unpredictability as evidenced by the prior art, and the amount of experimentation required to make and use all crystals and make and use all three-dimensional structures and methods of “rational drug design” as broadly encompassed by the claims, undue experimentation would be necessary for a skilled artisan to make and use the entire scope of the claimed invention.

Conclusion

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10. Status of the claims:

Claims 1-3, 7-9, 11, and 21-22 are pending.

Claims 1-3 and 7-8 appear to be in a condition for allowance.

Claims 9, 11, and 21-22 are rejected.


Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire **THREE MONTHS** from the mailing date of this action. In the event a first reply is filed within **TWO MONTHS** of the mailing date of this final action and the advisory action is not mailed until after the end of the **THREE-MONTH** shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than **SIX MONTHS** from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to David J. Steadman whose telephone number is 571-272-0942. The examiner can normally be reached on Mon to Fri, 7:30 am to 4:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Kathleen Kerr can be reached on 571-272-0931. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).


David J. Steadman, Ph.D.
Primary Examiner
Art Unit 1656

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APPENDIX A

CLUSTAL W (1.83) Multiple Sequence Alignments

Sequence type explicitly set to Protein

Sequence format is Pearson

Sequence 1: SEQ_ID_NO_3 435 aa
Sequence 2: SEQ_ID_NO_5 845 aa
Sequence 3: SEQ_ID_NO_6 708 aa
Sequence 4: SEQ_ID_NO_17 423 aa

Start of Pairwise alignments

Sequences (1:2) Aligned. Score: 100
Sequences (1:3) Aligned. Score: 100
Sequences (1:4) Aligned. Score: 100
Sequences (2:2) Aligned. Score: 100
Sequences (2:3) Aligned. Score: 69.9153
Sequences (2:4) Aligned. Score: 100
Sequences (3:2) Aligned. Score: 69.9153
Sequences (3:3) Aligned. Score: 100
Sequences (3:4) Aligned. Score: 100
Sequences (4:2) Aligned. Score: 100
Sequences (4:3) Aligned. Score: 100
Sequences (4:4) Aligned. Score: 100

Start of Multiple Alignment

SEQ_ID_NO_6 GLYSERHISMETSERPRVALPHETHRASNPASNSERSERPRPRALAVALPRGLNSERPHEG
SEQ_ID_NO_17 -----SERPRVALPHETHRASNPASNSERSERPRPRALAVALPRGLNSERPHEG
SEQ_ID_NO_3 GLYSERHISMETSERPRVALPHETHRASNPASNSERSERPRPRALAVALPRGLNSERPHEG
SEQ_ID_NO_5 GLYSERHISMETSERPRVALPHETHRASNPASNSERSERPRPRALAVALPRGLNSERPHEG

SEQ_ID_NO_6 LNVALALAHISLEUHSALAPRTHRGLYSERGLYLYSSERTHRLYSVALPRALAALATYR
SEQ_ID_NO_17 LNVALALAHISLEUHSALAPRTHRGLYSERGLYLYSSERTHRLYSVALPRALAALATYR
SEQ_ID_NO_3 LNVALALAHISLEUHSALAPRTHRGLYSERGLYLYSSERTHRLYSVALPRALAALATYR
SEQ_ID_NO_5 LNVALALAHISLEUHSALAPRTHRGLYSERGLYLYSSERTHRLYSVALPRALAALATYR

SEQ_ID_NO_6 ALAALAGLNGLYTYRLYSVALLEUVALLEUASNPRSERVALALAALATHRLEUGLYPHEG
SEQ_ID_NO_17 ALAALAGLNGLYTYRLYSVALLEUVALLEUASNPRSERVALALAALATHRLEUGLYPHEG
SEQ_ID_NO_3 ALAALAGLNGLYTYRLYSVALLEUVALLEUASNPRSERVALALAALATHRLEUGLYPHEG
SEQ_ID_NO_5 ALAALAGLNGLYTYRLYSVALLEUVALLEUASNPRSERVALALAALATHRLEUGLYPHEG

SEQ_ID_NO_6 LYALATYRMETSERLYSALAHISGLYVALASPPRASNILEARGTHRGlyVALARGTHRIL
SEQ_ID_NO_17 LYALATYRMETSERLYSALAHISGLYVALASPPRASNILEARGTHRGlyVALARGTHRIL
SEQ_ID_NO_3 LYALATYRMETSERLYSALAHISGLYVALASPPRASNILEARGTHRGlyVALARGTHRIL
SEQ_ID_NO_5 LYALATYRMETSERLYSALAHISGLYVALASPPRASNILEARGTHRGlyVALARGTHRIL

SEQ_ID_NO_6 ETHRTHRGlySERPRIETHRTYRSERTHRTYRGlyLYSPHELEUALAASPGlyGLYCYS
SEQ_ID_NO_17 ETHRTHRGlySERPRIETHRTYRSERTHRTYRGlyLYSPHELEUALAASPGlyGLYCYS
SEQ_ID_NO_3 ETHRTHRGlySERPRIETHRTYRSERTHRTYRGlyLYSPHELEUALAASPGlyGLYCYS
SEQ_ID_NO_5 ETHRTHRGlySERPRIETHRTYRSERTHRTYRGlyLYSPHELEUALAASPGlyGLYCYS

SEQ_ID_NO_6 SERGLYGLYALATYRASPILEILEILECYSASPGLUCYSHISSERTHRASPALATHRSER
SEQ_ID_NO_17 SERGLYGLYALATYRASPILEILEILECYSASPGLUCYSHISSERTHRASPALATHRSER
SEQ_ID_NO_3 SERGLYGLYALATYRASPILEILEILECYSASPGLUCYSHISSERTHRASPALATHRSER
SEQ_ID_NO_5 SERGLYGLYALATYRASPILEILEILECYSASPGLUCYSHISSERTHRASPALATHRSER

SEQ_ID_NO_6 ILELEUGLYILEGLYTHRVALLEUASPGNLALAGLUTHRALAGLYALAARGLEUVALVAL
SEQ_ID_NO_17 ILELEUGLYILEGLYTHRVALLEUASPGNLALAGLUTHRALAGLYALAARGLEUVALVAL
SEQ_ID_NO_3 ILELEUGLYILEGLYTHRVALLEUASPGNLALAGLUTHRALAGLYALAARGLEUVALVAL
SEQ_ID_NO_5 ILELEUGLYILEGLYTHRVALLEUASPGNLALAGLUTHRALAGLYALAARGLEUVALVAL

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SEQ_ID_NO_6 LEUALATHRALATHRRPRPRGLYSERGLYMETPHEASPSERSERVALLEU-----
SEQ_ID_NO_17 LEUALATHRALATHR-----
SEQ_ID_NO_3 LEUALATHRALATHR-----
SEQ_ID_NO_5 LEUALATHRALATHRRPRPRGLYSERVALTHRVALPRHISPRASNILEGLUGLUVALALAL

SEQ_ID_NO_6 -----CYSGLU--CYSTYRASPALAGLYCYSALATRPTYRG-----
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 EUSERTHRTHRGLYGLUILEPRPHETRYGLYLYSALAILEPRLEUGLUVALILELYSGLY

SEQ_ID_NO_6 -----LU-----LEUTHRPRALAGLUTHRTHR
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 GLYARGHISLEUILEPHECYSHISSERLYSLYSLYSCYSASPGLULEUALAALALYSLEU

SEQ_ID_NO_6 VALARGLEU-----ARGALATYRMETASNTHRPRGLYLEU-----PRV
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 VALALALEUGLYILEASNALAVALALATYRTRYRARGGLYLEUASPVALSERVALILEPRT

SEQ_ID_NO_6 ALCYSGNLNASPHISLEU-----GLUPHETRPGLU-----GLYVALPHETHRGLYLEU-
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 HRASNGLYASPVALVALVALVALALATHRASPALALEUMETTHRGLYPHETHRGLYASPP

SEQ_ID_NO_6 -----THRHSILEASPALAHISPHELEU-----SERGLNTH
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 HEASPSERVALILEASPCYSASNTHRSERASPGLYLYSPRGLNASPALAVALSERARGTH

SEQ_ID_NO_6 RLYSGNLNSERGLYGLUASNPHPEPTYRLEUVALALATYRGLNALATHRVALCYSALAARG
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 RGLNARGARGGLYARGTHRGLYARGGLYLYSPRGLYILETYRARGPHEVALALAPRGLYG

SEQ_ID_NO_6 ALAGLN
SEQ_ID_NO_17 -----
SEQ_ID_NO_3 -----
SEQ_ID_NO_5 LUARG-

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APPENDIX B

Seq1 is SEQ ID NO:1, Seq2 is SEQ ID NO:17

s-w opt: 2614 Z-score: 3193.8 bits: 602.0 E(): 4.3e-176
Smith-Waterman score: 2614; 100.000% identity (100.000% ungapped) in 412 aa overlap (510-921:1-412)

480	490	500	510	520	530
Seq1	EILEPRVALGLASNLEGLTHRTHRMETARGSERPRVALPHETHRASPNSESRERPRPR				
Seq2			SERPRVALPHETHRASPNSESRERPRPR		
			10	20	30

540	550	560	570	580	590
Seq1	ALAVALPRGLNLSERPHEGLNVALALAHISLEHISALAPRTHRGLYSERGLYLYSSERTHR				
Seq2	ALAVALPRGLNLSERPHEGLNVALALAHISLEHISALAPRTHRGLYSERGLYLYSSERTHR				
	40	50	60	70	80
					90

600	610	620	630	640	650
Seq1	LYSVALPRALAALATYRALAALAGLNGLYTYRLYSVALLEVALLEASNPRSERVALALAA				
Seq2	LYSVALPRALAALATYRALAALAGLNGLYTYRLYSVALLEVALLEASNPRSERVALALAA				
	100	110	120	130	140
					150

660	670	680	690	700	710
Seq1	LATHRLEGLYPHEGLYALATYRMETSERLYSALAHISGLYVALASPPRASNILEARGTHR				
Seq2	LATHRLEGLYPHEGLYALATYRMETSERLYSALAHISGLYVALASPPRASNILEARGTHR				
	160	170	180	190	200
					210

720	730	740	750	760	770
Seq1	GLYVALARGTHRILETHRTHRGLYSERPRILETHRTYRSERTHRTYRGLYLYSPHELEAL				
Seq2	GLYVALARGTHRILETHRTHRGLYSERPRILETHRTYRSERTHRTYRGLYLYSPHELEAL				
	220	230	240	250	260
					270

780	790	800	810	820	830
Seq1	AASPGLYGLYCYSSERGLYGLYALATYRASPILEILEILECYASPGLCYSHISSERTHR				
Seq2	AASPGLYGLYCYSSERGLYGLYALATYRASPILEILEILECYASPGLCYSHISSERTHR				
	280	290	300	310	320
					330

840	850	860	870	880	890
Seq1	ASPALATHRSERILELEGLYILEGLYTHRVALLEASPGLNALAGLTHRALAGLYALAARG				
Seq2	ASPALATHRSERILELEGLYILEGLYTHRVALLEASPGLNALAGLTHRALAGLYALAARG				
	340	350	360	370	380
					390

900	910	920	930	940	950
Seq1	LEVALVALLEALATHRALATHRPRPRGLYSERVALTHRVALPRHISPRASNILEGLGLVA				
Seq2	LEVALVALLEALATHRALATHR				
	400	410			

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APPENDIX C

```
>>Seq2                                     (820 aa)
  s-w opt: 4989  Z-score: 6113.3  bits: 1143.2  E():    0
Smith-Waterman score: 4989;  93.103% identity (98.901% ungapped) in 870 aa overlap (496-1364:1-820)
```

```

          470      480      490      500      510      520
Seq1  LYSALAVALASPPHEILEPRVALGLASNLEGLTHRTH-RMETARGSERPRVALPHETHRA
                               :: . : ::: : ::::::::::::::
Seq2                                GLYSERHISMET---SERPRVALPHETHRA
                                   .10                20

```

Seq1 SPASNSERSERPRPRALAVALPRLGNSERPHEGLNVALALAHISLEHISALAPRTHRGLY
 Seq2 SPASNSERSERPRPRALAVALPRLGNSERPHEGLNVALALAHISLEHISALAPRTHRGLY

```

          590      600      610      620      630      640
Seq1  SERGLYLYSSERTHRLYSVALPRALAALATYRALAALAGLNGLYTYRLYSVALLEVALLE
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
Seq2  SERGLYLYSSERTHRLYSVALPRALAALATYRALAALAGLNGLYTYRLYSVALLEVALLE
      90      100      110      120      130      140

```

```

      650      660      670      680      690      700
Seq1  ASNPRSERVALALAALATHRLEGLYPHEGLYALATYRMETSERLYSALAHISGLYVALAS
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Seq2  ASNPRSERVALALAALATHRLEGLYPHEGLYALATYRMETSERLYSALAHISGLYVALAS
      150      160      170      180      190      200

```

```

          710      720      730      740      750      760
Seq1  PPRASNILEARGTHRGLYVALARGTHRILETHRTHRGLYSERPRILETHRTYRSERTHRT
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
Seq2  PPRASNILEARGTHRGLYVALARGTHRILETHRTHRGLYSERPRILETHRTYRSERTHRT
      210      220      230      240      250      260

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Seq1 YRGLYLSPHELEALAASPGLYGLYCYSSERGLYGLYALATYRASPILEILEICYSAS
 Seq2 YRGLYLSPHELEALAASPGLYGLYCYSSERGLYGLYALATYRASPILEILEILEICYSAS

Seq1 PGLCYSHISSERTHRASPALATHRSERILEGLYILEGLYTHRVALLEASPLNALAGL

 Seq2 PGLCYSHISSERTHRASPALATHRSERILEGLYILEGLYTHRVALLEASPLNALAGL
 330 340 350 360 370 380

	890	900	910	920	930	940
Seq1	THRALAGLYALAARGLEVALVALL	EALATHRALATHRPRPGLYSERVA	LTHRVALPRHI	:	:	:
Seq2	THRALAGLYALAARGLEVALVALL	EALATHRALATHRPRPGLYSERVA	LTHRVALPRHI	:	:	:
	390	400	410	420	430	440

[illegible]

	1010	1020	1030	1040	1050	1060
Seq1	LEGLVALILELYSGLYGLYARGHISLEILEPHECYSHISSERLYSLYSLYSCYSPG	L				
	:	:	:	:	:	:
Seq2	LEGLVALILELYSGLYGLYARGHISLEILEPHECYSHISSERLYSLYSLYSCYSPG	L				
	510	520	530	540	550	560

	1070	1080	1090	1100	1110	1120
Seq1	EALAAALYSLEVALALALEGLYILEASNALAVALLATYRTRYRAGGLYLEASPVALSE					
Seq2	EALAAALYSLEVALALALEGLYILEASNALAVALLATYRTRYRAGGLYLEASPVALSE					
	570	580	590	600	610	620
	1130	1140	1150	1160	1170	1180
Seq1	RVALILEPRTHRASNGLYASPVALLVALVALALATHRASPALALEMETTHRGlyPHET					
Seq2	RVALILEPRTHRASNGLYASPVALLVALVALALATHRASPALALEMETTHRGlyPHET					
	630	640	650	660	670	680
	1190	1200	1210	1220	1230	1240
Seq1	HRGLYASPPHEASPSERVALILEASPCYSASNTHRCYSVALTHRGLNTHRVALASPPHES					
Seq2	HRGLYASPPHEASPSERVALILEASPCYSASNTHR-----S					
	690	700	710	720		
	1250	1260	1270	1280	1290	1300
Seq1	ERLEASPPRTHRPHETHRILEGLTHRTHRTHRLERPGNLNAPALAVALLSERARGTHRGLN					
Seq2	ER--ASP-----GL-----YLYSPRGNLNASPALAVALLSERARGTHRGLN					
			730	740	750	760
	1310	1320	1330	1340	1350	1360
Seq1	ARGARGGLYARGTHRGLYARGGGLYLYSPRGLYILETYRARGPHEVALALAPRGLYGLARG					
Seq2	ARGARGGLYARGTHRGLYARGGGLYLYSPRGLYILETYRARGPHEVALALAPRGLYGLARG					
	770	780	790	800	810	820
	1370	1380	1390	1400	1410	1420
Seq1	PRSERGLYMETPHEASPSERSEVALLECYSGLCYSTYRASPALAGLYCYSALATRPTRY					